

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
7 November 2002 (07.11.2002)

PCT

(10) International Publication Number
WO 02/087731 A1(51) International Patent Classification⁷: **B01D 53/32.**
B01J 19/08, F01N 3/08LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(21) International Application Number: PCT/GB02/01820

(22) International Filing Date: 19 April 2002 (19.04.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0110345.6 27 April 2001 (27.04.2001) GB

(71) Applicant (for all designated States except US): ACCENTUS PLC [GB/GB]; 329 Harwell, Didcot, Oxfordshire OX11 0QJ (GB).

(72) Inventors; and

(75) Inventors/Applicants (for US only): MORGAN, Ross, Alexander [GB/GB]; 28 Buckingham Street, Grandpont, Oxford, Oxfordshire OX1 4LH (GB). SEGAL, David, Leslie [GB/GB]; 43 Foxborough Road, Radley, Abingdon, Oxfordshire OX14 3AB (GB). SHAWCROSS, James, Timothy [GB/GB]; 27 Ticknell Piece Road, Charlbury, Oxfordshire OX7 3TN (GB).

(74) Agents: LOFTING, Marcus, John et al.; c/o Accentus plc, Patents Dept., 329 Harwell, Didcot, Oxfordshire OX11 0QJ (GB).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, I.K, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW. ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

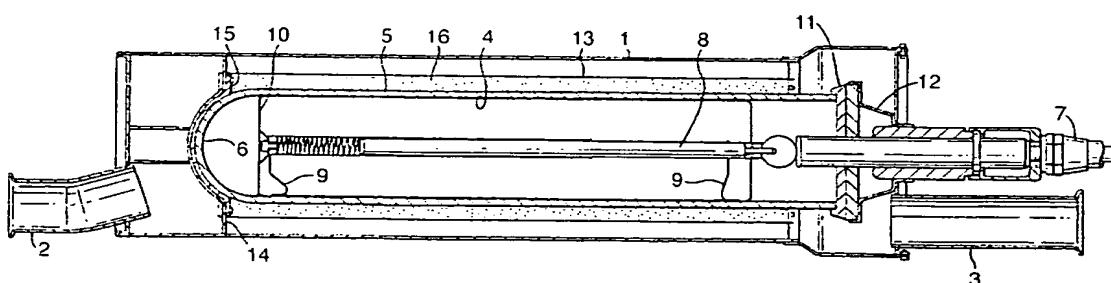
- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, I.K, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW. ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)
- of inventorship (Rule 4.17(iv)) for US only

Published:

- with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: REACTOR FOR TRAPPING AND OXIDATION OF CARBONACEOUS MATERIAL



(57) Abstract: A reactor (1) comprises a pair of electrodes (4, 13) spaced apart to provide a path therebetween for gas flow. A permeable mass of silicon carbide (16) is provided in the space between the electrodes (4, 13) for trapping carbonaceous material in the gas flowing through the space. The silicon carbide (16) is arranged to have an electrical resistivity which is sufficiently high to permit formation, when an appropriate electrical potential is applied across the electrodes (4, 13), of a plasma in the gas within the interstices of the silicon carbide through which the gas permeates.

WO 02/087731 A1

- 1 -

Reactor for trapping and oxidation of carbonaceous material.

The invention relates to a reactor for the trapping
5 and removal of carbonaceous material in a gas stream such
as the exhaust of an internal combustion engine.

A number of approaches are currently used for the removal of carbonaceous material and other combustion
10 products from the exhaust of internal combustion engines and diesel engines in particular. The term carbonaceous material refers to carbonaceous particulates commonly referred to as soot, hydrocarbons including polycyclic aromatic hydrocarbons and soluble organic fractions resulting from
15 fuel combustion, carbon monoxide and other regulated or unregulated combustion products from the exhausts of internal combustion engines. Cordierite ceramic honeycombs that have alternate channels blocked have been evaluated and used for nearly two decades as diesel
20 particulate filters (DPF) and an early discussion of them is given by Wade et al in 'Diesel Particulate Trap Regeneration Techniques' published in SAE 810118 (1981). More recently Cutler and Merkel in 'A New High
Temperature Ceramic Material for Diesel Particulate
25 Filter Applications' in SAE 2000-01-2844, pages 79-87 have reviewed the use of cordierite-based diesel particulate filters. They refer to the geometric properties of a representative filter that is 5.66 inches in diameter, 6 inches in length containing 178 cells per
30 square inch with a wall thickness of 0.012 inches. The ends of alternate cells are blocked off so that exhaust gases pass through the porous walls between the cells and carbonaceous particulate material is trapped within the porous wall structure. Filters with alternate channels
35 blocked off are often referred to as wallflow filters. They can be regenerated by external heating to burn off trapped material or by use of a carbon combustion catalyst such as Eolys, a cerium oxide additive for fuel

- 2 -

described in WO 00/43102. Cordierite has a melting point of around 1460°C and can react with residues of metallic salt additives in lubricating oil and diesel fuel when high temperature excursions are experienced in the
5 exhaust gases to form tightly bonded glass-like deposits on the filter. The low thermal conductivity of cordierite ($<2 \text{ W m}^{-1} \text{ K}^{-1}$) makes dissipation of heat by the filter material more difficult and can encourage formation of these glass-like deposits. It is difficult
10 to regenerate the filter once the glass is produced and in addition melting and deformation of the cordierite filter can occur at high temperature.

Silicon carbide has traditionally found application
15 as heating elements as highlighted by Moulson and Herbert in 'Electroceramics: Materials: Properties: Applications', published by Chapman & Hall, 1993, page 121 who refer to this use that is aided by the low electrical resistivity of this ceramic material. Use of
20 silicon carbide as a substrate for the treatment of vehicle emissions is a recent development and has been restricted to non-plasma aftertreatment processes.

In order to avoid the above problems arising from
25 the use of cordierite DPF, silicon carbide filters of a similar geometry to conventional diesel particulate filters have been used as discussed by Ohno et al in 'Characterization of SiC-DPF for Passenger Car' in SAE 2000-01-0185 pages 25-38. Silicon carbide can operate at
30 temperatures higher than cordierite as its sublimation temperature 2200°C is much higher than the melting point of cordierite, 1460°C. Silicon carbide has a higher thermal conductivity than cordierite, $73 \text{ W m}^{-1} \text{ K}^{-1}$ compared to $2 \text{ W m}^{-1} \text{ K}^{-1}$ according to Ohno et al. These properties
35 together with the lower chemical reactivity of silicon carbide towards additives in the lubricating oil and diesel fuel considerably reduce the tendency to form glass-like deposits on silicon carbide filters. Silicon

carbide filters can be regenerated by the same methods used for regeneration of cordierite filters. Morawietz in 'Not Just Clean, but Pure' in Automotive Business International January/February 2001, pages 68-70
5 describes the use of a silicon carbide filter in combination with a cerium-based carbon combustion catalyst.

Silicon carbide fibre has also been used for
10 construction of diesel particulate filters for removing carbonaceous particulate material. EP 0 742 352 B (Isuzu) describes a diesel particulate filter consisting of long silicon carbide fibre that can be in the form of a felt. The filter is regenerated by electrically heating a
15 conducting metallic net that heats up the fibre and burns off particulate material. The electrical resistivity of silicon carbide has not been referred to in the above examples as an important parameter for regeneration of the diesel particulate filter. Continuous silicon carbide
20 fibre, in contrast to whiskers or chopped fibre can be prepared by a process known as polymer pyrolysis from polymers known as polycarbosilanes as described by Riedel in 'Advanced Ceramics from Inorganic Polymers', in Processing of Ceramics, edited by RJ Brook, VCH
25 Publishers, (1996), pages 1-50. The pyrolysis of polycarbosilanes has led to the commercial production of continuous silicon carbide fibre known as Nicalon™ (Nippon Carbon) and Tyranno™ (Ube Industries). It is possible to obtain continuous silicon carbide fibre with
30 a range of electrical resistivities. For example, high volume resistivity grade Nicalon™ has a volume resistivity of 10^6 ohm cm while low volume resistivity grade Nicalon™ has a resistivity of 0.5 - 5.0 ohm cm as described in the Technical Leaflet: Nicalon™ Fiber
35 published by Nippon Carbon Company Limited (1998). The technical leaflet Polycarbosilane 'Nipusi®' published by Nippon Carbon Company (1995) indicates that the atmosphere in which polymer pyrolysis takes place,

- 4 -

oxidising or inert, can affect the electrical resistivity of the resulting silicon carbide fibre.

None of the above uses of silicon carbide for the treatment of vehicle exhaust emissions involve the use of non-thermal plasmas. There is increasing interest in the use of non-thermal plasmas for treatment of gaseous exhausts, in particular for treatment of exhausts from motor vehicles. Examples of non-thermal plasma treatment of exhausts are described in US 3,983,021 (Monsanto), US 5,147,516 (Tecogen) and US 5,254,231 (Battelle Memorial Institute). GB 2,274,412 (AEA Technology) describes a method for the treatment of diesel emissions by a non-thermal plasma for oxidation of carbonaceous particulates and reduction of NO_x to nitrogen. Plasma can be used to activate or produce reactant species, which then subsequently react with or without catalytic enhancement to yield the desirable products. For example, WO99/12638 describes the plasma production of plasma activated hydrocarbons as a precursor to the selective catalytic reduction of NO_x to N₂. Other examples of plasma and plasma-catalyst systems for treatment of exhaust gases are described in WO 00/29727 (Engelhard), US 6038854 (Regents of the University of California) and WO 00/21646 (Johnson Matthey).

There is a requirement for a material for treatment of carbonaceous particulate in exhaust gases, particularly in diesel exhaust gases from vehicles, which avoids the disadvantages of cordierite and that can be used in a non-thermal plasma. The low electrical resistivity of silicon carbide renders it unsuitable for use as a trap for carbonaceous matter in a plasma reactor because the low electrical resistivity prevents establishment of the required electric field across the trap for generation of the plasma.

- 5 -

It is an object of the invention to provide a reactor incorporating a trapping material for carbonaceous matter which avoids the disadvantages of cordierite and is also suitable for use in a non-thermal 5 plasma.

The invention is not restricted to a particular type of non-thermal reactor and may be put into practice in dielectric barrier reactors also known as silent 10 discharge reactors, packed bed reactor also known as pellet bed reactor, pulsed corona reactor, microwave reactor or surface discharge reactor.

The invention provides a reactor for the trapping 15 and oxidation of carbonaceous material in a gas stream, such as the exhaust from an internal combustion engine, which reactor comprises a pair of electrodes spaced apart and adapted on application of a suitable electrical potential thereacross for generating a plasma in gas 20 flowing through the space between the electrodes, characterised in that a gas permeable mass of silicon carbide is provided in the space between the electrodes for trapping carbonaceous material thereon, and in that provision is made for the mass of silicon carbide to have 25 an electrical resistivity which is sufficiently high to permit formation of a plasma in gas within the interstices of the silicon carbide through which the gas permeates.

30 Preferably the reactor is a dielectric barrier reactor, there being provided a continuous dielectric layer between the said pair of electrodes.

Also, in operation of a reactor according to this 35 invention, the trapped material can be activated by the plasma or trapping material to a state where it can react with for example NO_x to yield desirable products. The permeable mass of silicon carbide, or a coating or

- 6 -

constituent thereof, or the trapped species itself may, in the presence of a plasma act as a catalytic surface but importantly neither the plasma nor the permeable mass nor the trapped species alone need necessarily have
5 intrinsic catalytic properties. One example of a trapped species is carbonaceous particulate material from a diesel engine, for example soot that consists mainly of elemental carbon. In the plasma region soot becomes exposed to plasma generated species for example oxygen
10 atoms. Oxygen atoms or other plasma generated species may diffuse into, adsorb and react with soot. Other plasma generated species include but are not restricted to OH, O₃ and NO₂. For example it is known oxygen atoms can diffuse into soot and form aldehyde-type groups on
15 the surface. Oxygenated soot has different activated and catalytic properties to non-oxygenated soot.

The gas permeable mass of silicon carbide may be provided in the form of a monolith, wall flow honeycomb
20 filter, foam, graded foam, plates, fibres, meshes or weaves or combinations of these shapes. Fibres can be in the form of continuous or chopped fibre, felt, mat or blanket where the fibres can be randomly or non-randomly oriented. The required high electrical
25 resistivity of the silicon carbide is preferably provided by doping of the silicon carbide during manufacture or by a heat treatment of the silicon carbide in a controlled gaseous atmosphere, the doping and heat treatment being of an extent to allow the establishment of an electric
30 field and plasma across the material. For example a silicon carbide identified as SC-211 (<http://www.kyocera.com>) is commercially available from Kyocera which has a volume electrical resistivity of 8x10¹⁴ ohm cm at ambient temperature. This grade of
35 silicon carbide can be fabricated as a powder and the powder formed into a number of shapes such as plates, discs and cylinders.

- 7 -

Our investigations have indicated that a volume electrical resistivity of 10^6 ohm cm at ambient temperature or greater is required if non-thermal plasma is to be satisfactorily established within the 5 interstices of the silicon carbide through which the gas permeates. Preferably the volume electrical resistivity is significantly greater than 10^6 ohm cm.

For silicon carbide in the form of fibres, it is 10 possible for the required high electrical resistivity to be provided by the gaseous atmosphere used when polymer pyrolysis is carried out during manufacture. High volume resistivity grade Nicalon™ has a volume resistivity of 10^6 ohm cm and can be used. This value of resistivity is the 15 maximum available commercially for the fibre. However a post manufacturing treatment may be used to alter the electrical resistivity of silicon carbide fibres. For example an adherent ceramic coating can be deposited onto the fibres by a gas phase reaction for example by 20 chemical vapour deposition. Silica coatings may be suitable for this purpose.

Specific constructions of reactor embodying the invention will now be described by way of example and 25 with reference to the drawings filed herewith in which:-

Figure 1 is a longitudinal section of a reactor embodying the invention for the removal of carbonaceous combustion products and other combustion products from 30 the exhaust emissions of internal combustion engines and

Figure 2 is a schematic view showing the gas path through the embodiment shown in Figure 1. Both Figure 1 and Figure 2 are described in application WO 00/71866.

35

Referring to Figure 1 a reactor for the plasma assisted processing of the exhaust emissions from internal combustion engines to remove noxious components

therefrom consists of a reactor chamber 1 which has inlet and outlet stubs 2, 3, respectively by means of which it can be incorporated into the exhaust system of an internal combustion engine.

5

Inside the reactor chamber 1 there is a an inner electrode 4 which is supported within a dielectric tube 5 made for example out of α -alumina which has its upstream end closed by a spherical dome 6 to facilitate the flow 10 of exhaust gases through the reactor. The inner surface of the dielectric tube 5 can be metallised with a metal coating in order to increase the physical contact between the electrode and dielectric tube although in this example, the inner electrode 4 is conveniently provided 15 by a deposited electrically conducting layer of silver on the inner surface of the dielectric tube 5. High voltage connection via a high voltage input terminal 7 is made through a spring loaded telescopic tube assembly 8 and spring contacts 9. Load from the sprung telescopic tube 20 assembly 8 is received by a load spreader plate 10, which is connected to the conducting layer of silver forming the inner electrode 4. The materials including the spring are required to operate at elevated temperatures and the spring must have low creep at such temperatures. 25 A preferred material for the spring is an Inconel alloy such as X750. Alumina end flange 11 is shaped to receive and locate the end of the dielectric tube 5 and is itself located by a sprung metal clip 12. The space between the dielectric tube 5, on the inner surface of which is the 30 inner electrode 4, and the outer electrode 13 is filled with a gas permeable mass of silicon carbide illustrated diagrammatically at 16. This mass of silicon carbide may be in the form of a foam, grade foam, wallflow honeycomb filter, plates, porous particles, fibres, meshes, weaves 35 or a self-supporting hollow cylinder constructed of silicon carbide with a fine pore structure so that it is gas permeable or combinations of these shapes.

- 9 -

A potential of the order of kilovolts to tens of kilovolts and repetition frequencies in the range 50 to 5000 Hz can be applied to the inner electrode 4 through the high voltage input terminal 7. Pulsed direct current
5 is convenient for automotive use but alternating potentials for example triangular or sine waves or square wave or saw tooth wave of the same or similar characteristics can be used separately or in combination.
Concentric with the inner electrode 4 and dielectric
10 tube 5 is a grounded outer electrode 13 made for example of stainless steel. At the inlet end of the reactor the spherical dome of the dielectric tube 5 is in contact with a compliant heat resistant material 14 that rests in the curved part of the outer electrode 13 and held in
15 place by a metallic ring 15 with a series of screws.

As shown in Figure 2, the outer electrode 13 has a series of baffles 216 and slots 217, 217a. The baffles 216 extend from the outer electrode 213 to the inner
20 surface of the wall of the reactor chamber 21 and act as grounding connections as well as causing the exhaust gases to follow a convoluted path which has both axial, and circumferential components and being at least partially helical. There is also a radial component of
25 flow, initially inwardly as the gas transfers from outside the outer electrode 213 to the space between the dielectric tube 5 and the outer electrode 13 and then outwardly as the gas returns to exit from outside the outer electrode 13. Thus there is also a spiral component
30 in the flow.

The baffle 216 is arranged to divide the space between the electrode 13 and the reactor chamber 1 into six segments. At the gas inlet end three of these
35 segments are closed off at 216a, 216b and 216c to axial gas flow and the remaining three segments are open to axial gas flow into the space between the electrode 13 and the reactor chamber 1. These latter three segments

- 10 -

are closed off by the baffle 216 at the gas outlet end of the reactor. Consequently the gas is forced to pass via slot 217 radially into the space between the electrodes 13 and 4 then passing in at least a partially helical 5 manner before passing radially via the next slot 217a into the next segment of space between electrode 13 and reactor chamber 1. The baffle 216 leaves open this segment at the gas outlet end, allowing exhaust of the treated gas to exit the reactor. Thus it will be seen 10 that the exhaust gases both enter and leave the main part of the reactor 1 along the surface of the outer electrode 13. Thus for a given gas velocity, the residence time of the exhaust gases in the electric field is increased compared with either purely axial or radial flow. Note 15 that in Figure 2 part of the electrode 13 has been shown cut away at 218. This cut away is shown in Figure 2 only to illustrate the flow of the exhaust gases as they pass between the electrodes and does not represent a structural feature of the reactor.

20

In operation, the exhaust gas is caused to permeate through the mass of silicon carbide 16. In so doing, carbonaceous particulates and other carbonaceous matter become trapped on the silicon carbide. Non-thermal 25 plasma, sustained in the gas in the interstices of the gas permeable mass of silicon carbide 16, by the applied electrical potential, is effective both directly and via active species produced in the plasma to oxidise the trapped carbonaceous matter.

30

When the silicon carbide 16 is in the form of a wallflow honeycomb it can consist of an array of channels, typically 200 per square inch with alternate channels blocked off at their ends, extending along the 35 length of the honeycomb parallel with the axis of the honeycomb although other forms of silicon carbide can be used or a packing of silicon carbide fibres or porous particles. For example a foam or other suitable

- 11 -

monolithic form can be used or a packing of silicon carbide fibres or porous particles. If a foam is used a sintering aid to aid diffusion or encourage liquid-phase sintering during manufacture, for example aluminium oxide, may be present in the silicon carbide as silicon carbide by itself is particularly difficult to shape by sintering because of the essentially covalent nature of this material. Reaction bonded silicon carbide shapes can also be used and can be produced by infiltration of molten silicon into a ceramic body shaped out of α -silicon carbide powder and a carbon source, a powder or phenolic resin as examples. Such shapes do not exhibit significant dimensional changes on reaction with molten silicon. Excess porosity in the reaction bonded silicon carbide is filled with silicon during the reaction bonding process. An example of reaction bonded silicon carbide is REFELTM (British Nuclear Fuels).

In all cases, however, it is important that the silicon carbide has high electrical resistivity, as explained so that non-thermal plasma can be produced in gas within the interstices of the silicon carbide by application of the electrical potential. This can be provided inherently in the silicon carbide itself by adopting material commercially available as SC-211 from Kyocera or commercially available fibres with a suitable electrical resistivity. Our investigations have indicated that the volume electrical resistivity should be at least 10^6 ohm cm and preferably significantly higher. If fibres are used, electrical resistivity can be provided by a suitable coating applied to the fibres subsequently to their manufacture. In this case silicon carbide fibres available from Nippon Carbon as NicalonTM or from Ube Industries as TyrannoTM can be used. Both of these are derived from polycarbosilane precursors, while Tyranno also incorporates titanium or zirconium into the fibre structure.

- 12 -

Operation of the reactor can be further enhanced by provision of a catalytic coating on the silicon carbide.

Such a catalytic coating may enhance the oxidation process of trapped carbonaceous matter and/or catalyse

5 other desired reactions on other noxious constituents of the permeating gas such as the catalytic removal of nitrogeneous oxides. Suitable catalytic materials are described in WO 99/12638, WO 00/43102, WO 99/38603 and 00 15952.5 (UK filed 30 June 2000).

10

The invention is not restricted to the details of the foregoing examples. For instance, other configurations of reactor can be used, although the particular form in which the mass of gas permeable 15 silicon carbide is provided has to be selected according to the nature of the gas flow path provided. Subject to this, examples of other reactor configurations which may be used are described in patent publications WO 00/71866, WO 99/47243, WO 99/67510 or WO 99/12638 as well as other 20 types of non-thermal plasma reactors such as pulsed corona reactors and surface discharge reactors. When alternating potentials are used triangular waves, sine waves, square waves or saw-tooth waves can be used separately or in combination as can pulsed direct current 25 for other reactor configurations.

- 13 -

Claims.

1. A reactor for the trapping and oxidation of carbonaceous material in a gas stream, which reactor comprises a pair of electrodes (4, 13) spaced apart and
5 adapted on application of a suitable electrical potential thereacross for generating a plasma in gas flowing through the space between the electrodes (4, 13), characterised in that a gas permeable mass of silicon carbide (16) is provided in the space between the
10 electrodes for trapping carbonaceous material thereon, and in that provision is made for the mass of silicon carbide (16) to have an electrical resistivity which is sufficiently high to permit formation of a plasma in gas within the interstices of the silicon carbide (16)
15 through which the gas permeates.

2. A reactor as claimed in claim 1, in which there is provided a continuous dielectric layer (5) between the said pair of electrodes (4, 13), whereby the reactor is a
20 dielectric barrier reactor.

3. A reactor as claimed in claim 1 or claim 2, in which the gas permeable mass of silicon carbide (16) is
25 provided in the form of a monolith, honeycomb, wall flow filter, foam, plates or fibres.

4. A reactor as claimed in claim 3, further characterised in that the gas permeable mass of silicon
30 carbide (16) is in the form of fibres and the high electrical resistivity is provided by a coating on the fibres.

5. A reactor as claimed in claim 3, further
35 characterised in that the gas permeable mass of silicon carbide (16) is so made as to possess in bulk the said sufficiently high electrical resistivity.

- 14 -

6. A reactor as claimed in claim 5, further characterised in that the volume electrical resistivity of the gas permeable mass of silicon carbide (16) is at least 10^6 ohm cm.

5

7. A reactor as claimed in claim 6, further characterised in that the gas permeable mass of silicon carbide (16) is fabricated from that designated as SC-211 from Kyocera having a volume electrical resistivity of
10 8×10^{14} ohm cm.

8. A reactor as claimed in any of the preceding claims, in which exposed surfaces of the gas permeable mass of silicon carbide are coated with a catalytic
15 material.

1/1

Fig.1.

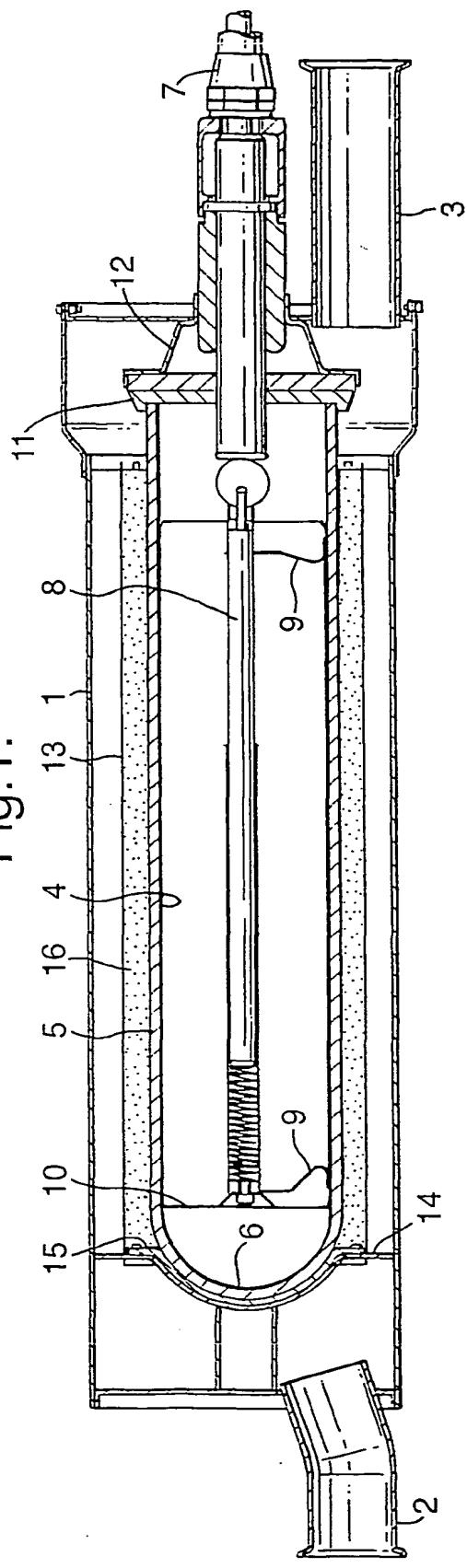
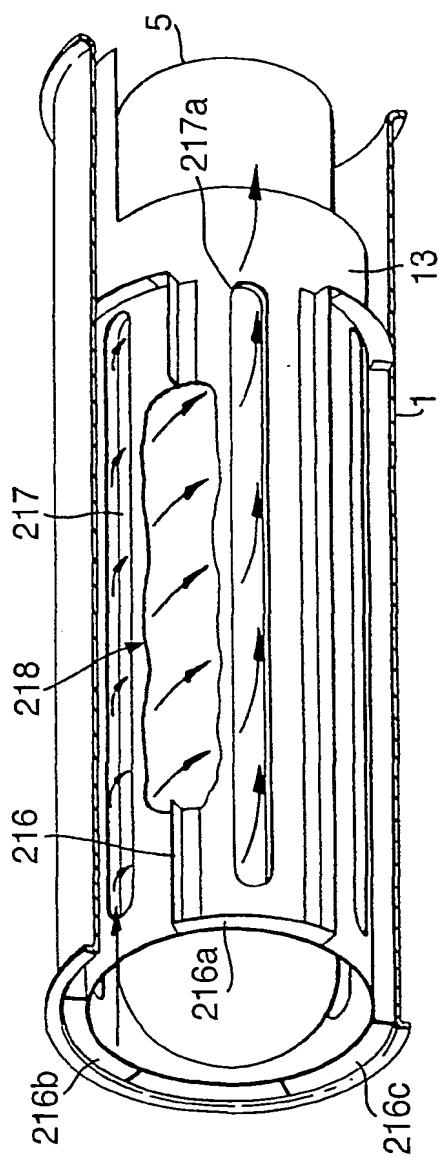


Fig.2.



INTERNATIONAL SEARCH REPORT

Intern

Application No

PC1/GB 02/01820

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 B01D53/32 B01J19/08 F01N3/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 7 B01J B01D F01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 01 30485 A (MARTIN ANTHONY ROBERT ; SHAWCROSS JAMES TIMOTHY (GB); RAYBONE DAVID) 3 May 2001 (2001-05-03) page 10, line 12-24; figures ---	1-3, 5, 6
P, X	DE 100 57 862 C (SIEMENS AG) 7 February 2002 (2002-02-07) column 7, paragraph 42 column 8, paragraphs 49,50; claims 15-27; figure 13 ---	1-3, 5, 6
A	DE 197 17 889 C (INST NIEDERTEMPERATUR PLASMAPH) 8 April 1999 (1999-04-08) column 4, line 4-8 ---	8
A	US 5 767 470 A (CHA CHANG YUL) 16 June 1998 (1998-06-16) the whole document ---	
	-/-	

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

17 July 2002

Date of mailing of the international search report

25/07/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Gruber, M

INTERNATIONAL SEARCH REPORT

Int'l Application No
PCT/GB 02/01820

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	EP 1 125 704 A (NGK INSULATORS LTD) 22 August 2001 (2001-08-22) column 20, paragraph 84 column 26, paragraph 105; claims 1,10; figure 21 -----	
P,A	WO 01 36805 A (SOUTHWEST RES INST) 25 May 2001 (2001-05-25) page 5, line 12-23; figure 2 -----	

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte.....	Application No
PCT/GB 02/01820	

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 0130485	A	03-05-2001	AU BR EP WO	7805300 A 0014900 A 1222014 A1 0130485 A1		08-05-2001 11-06-2002 17-07-2002 03-05-2001
DE 10057862	C	07-02-2002	DE WO	10057862 C1 0242615 A1		07-02-2002 30-05-2002
DE 19717889	C	08-04-1999	DE WO EP JP	19717889 C1 9843922 A1 0979135 A1 2001522302 T		08-04-1999 05-11-1998 16-02-2000 13-11-2001
US 5767470	A	16-06-1998	US	6027698 A		22-02-2000
EP 1125704	A	22-08-2001	EP CN WO	1125704 A1 1327410 T 0115877 A1		22-08-2001 19-12-2001 08-03-2001
WO 0136805	A	25-05-2001	US AU WO	2001042372 A1 2116801 A 0136805 A1		22-11-2001 30-05-2001 25-05-2001

THIS PAGE BLANK (USPTO)